U.S. Patent and Trademark Office, U.S. DEPARTM Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless of contains a valid

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10715741		
Filing Date		2003-11-18		
First Named Inventor	Kenji Kimura			
Art Unit		2624		
Examiner Name				
Attorney Docket Number	or	DR414a		

					U.S.	PATENTS			Remove		
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue E	Date	Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear			
	1										
If you wisl	h to a	l dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.		Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	of sited Desument Rele			ges,Columns,Lines where levant Passages or Relevant ures Appear		
	1										
If you wis	h to a	dd additional U.S. Publi		p		, , , , , , , , , , , , , , , , , , , ,		d button			
				FOREIG	SN PAT	TENT DOCUM	ENTS		Remove		
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or	Pages,Colum where Releva Passages or l Figures Appe	nt Relevant	70
	1	3177156	JP			1991-08-01	CANON INC				V
	2	61111063	JP			1986-05-29	CANON INC				
	3	01258557	JP			1989-10-16	RICOH CO LTD				

	4	04306057	JP		1992-10-28	MITSUBISHI ELECTRIC CORP		
	5	10107962	JP		1998-04-24	NEC CORP		Z
If you wis	h to a	dd additional Foreign F	atent Document	citation	information pl	lease click the Add butto	Add	_
			NON-PATE	NT LITE	ERATURE DO	CUMENTS	Remove	
Examiner Initials*	caminar Citie Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (blook, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city andior country where published.							Тs
	1							
If you wis	h to a	dd additional non-pater	nt literature docu	ment ci	tation informat	ion please click the Add	button Add	_
			EX	AMINE	R SIGNATUR	E		
Examiner	Signa	ture				Date Considered		
						ormance with MPEP 609 with next communication		

See Kind Code of USPTO Petent Documents at twent_ISETO_GOL/or MPEP 901.04. * Enter office that issued the document, by the two-lets of (WPD) Standard 513.) * For implanees petent focuments, the includance of the pear of the register or twent precede the serial number of the petent document. A produced the period of the petent of the pet

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10715741		
Filing Date		2003-11-18		
First Named Inventor	Kenji Kimura			
Art Unit		2624		
Examiner Name				
Attorney Docket Number		P8414a		

CERTIFICATION STATEMENT

Please see 3	7 CFR 1	.97 and	1.98 to make	the appropriate selection	on(s):
--------------	---------	---------	--------------	---------------------------	--------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 3.7 CFR 1.97(eVI.)

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.5(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.57(c) for the contraction of th

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- 7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Daniel A. Ratoff/	Date (YYYY-MM-DD)	2006-05-12
Name/Print	Daniel A. Ratoff	Registration Number	54389

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is foll field and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR 1.14. This collection is estimated to take it hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenan's Office, and Superiment of Commence, P. 0. Bot 1450, Alexandria, V.32.511.450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P. 0. Box 1450, Alexandria, V.32.313-1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.